AMENDMENTS

IN THE CLAIMS:

- 1. (Cancelled).
- 2. (Cancelled).
- 3. (Currently amended) The A method according to claim 2 of evaluating the efficiency of a sterilization process, which comprises the steps of:
 - a) <u>subjecting a sufficient amount of at least one prion protein degradation indicator in a</u>
 <u>container to said sterilization process; and</u>
 - b) determining the level of degradation of said indicator,
 - wherein said indicator of step a) is transcribed by a gene naturally occurring in a wherein said_fungus selected from the group consisting of Saccharomyces cerevisiae, and Podospora anserina.
- 4. (Original) The method according to claim 3, wherein said indicator is transcribed by a gene selected from the group consisting of SUP35, URE2 and HET-s.
- 5. (Currently amended) The method according to claim 2 3, wherein said indicator is selected from the group consisting of Sup35p, Ure2p, Het-s protein, and combination thereof.
- 6. (Currently amended) The method according to claim 1 3, wherein said indicator is a purified form naturally occurring in *Saccharomyces cerevisiae*, *Podospora anserina* or a fungus, a recombinant form, an analog, a mutant, or a fragment of said indicator.
- 7. (Currently amended) The method according to claim 4 3, wherein said indicator is a biological indicator, a biochemical indicator, or a chemical indicator.
- 8. (Currently amended) The method according to claim 4 3, wherein step b) is performed by determining the weight or the mass, quantifying radicals, colorimetric variations, radiometry, nephelometry, immuno-enzymatic method, Westerm blotting, dot blotting, radioimmuno

- assay, circular dichroism, electron microscopy, fluorescent microscopy, FTIR, Congo red binding, or proteinase digestion.
- 9. (Currently amended) The method according to claim ± 3 , wherein said sterilization process is performed by autoclaving, chemical exposure, dry heating, low temperature plasma gas, ozone-based exposure, or sterilization techniques using alkylant alkylating and/or oxidizing sterilizing agents.
- 10. (Currently amended) The method according to claim 4 <u>3</u>, wherein said chemical exposure is a vapor or a solution selected from the group consisting of detergent, ethylene oxide, protease, sodium hydroxide, and enzyme.
- 11. (Currently amended) The method of claim 1 3, wherein said amount of indicator of step a) is between 0.1 ng to 100 g.
- 12. (Currently amended) The method of claim 4 3, wherein said container is of a material selected from the group consisting of paper, glass, borosilicate, metal, polymer, alloy, and composite.
- 13. (Currently amended) The method according to claim 44 <u>3</u>, wherein said container is porous, permeable, or semi-permeable.